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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/019,785	01/04/2002	Hidemi Saito	04853.0087	1823
22852	7590 07/01/2004		EXAMINER	
FINNEGAN, HENDERSON, FARABOW, GARRETT & DUNNER			GALVEZ, JAMES JASON	
LLP 1300 I STRE	ET, NW		ART UNIT	PAPER NUMBER
WASHINGTON, DC 20005			1647	
			DATE MAILED: 07/01/2004	4

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)	
	10/019,785	SAITO ET AL.	
Office Action Summary	Examiner	Art Unit	
	J. Jason Galvez	1647	
The MAILING DATE of this communication Period for Reply	appears on the cover sheet with	n the correspondence address	
A SHORTENED STATUTORY PERIOD FOR RETHE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CF after SIX (6) MONTHS from the mailing date of this communication. If the period for reply specified above is less than thirty (30) days, If NO period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by some Any reply received by the Office later than three months after the rearned patent term adjustment. See 37 CFR 1.704(b).	ON. FR 1.136(a). In no event, however, may a reply. In a reply within the statutory minimum of thirty eriod will apply and will expire SIX (6) MONT statute, cause the application to become ABA	oly be timely filed (30) days will be considered timely. HS from the mailing date of this communication. NDONED (35 U.S.C. § 133).	
Status			
Responsive to communication(s) filed on _ This action is FINAL . 2b) Since this application is in condition for all closed in accordance with the practice und	This action is non-final. owance except for formal matte	· •	
Disposition of Claims			
4) ⊠ Claim(s) <u>1-15</u> is/are pending in the applica 4a) Of the above claim(s) is/are with 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) <u>1-15</u> are subject to restriction and	ndrawn from consideration.		
Application Papers			1
9) The specification is objected to by the Exar 10) The drawing(s) filed on is/are: a) Applicant may not request that any objection to Replacement drawing sheet(s) including the co	accepted or b) objected to be the drawing(s) be held in abeyand prection is required if the drawing(s	e. See 37 CFR 1.85(a).) is objected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. § 119			
12) Acknowledgment is made of a claim for force a) All b) Some * c) None of: 1. Certified copies of the priority docum 2. Certified copies of the priority docum 3. Copies of the certified copies of the application from the International Bu * See the attached detailed Office action for a	nents have been received. nents have been received in Ap priority documents have been r ireau (PCT Rule 17.2(a)).	plication No eceived in this National Stage	
Attachment(c)			
Attachment(s) 1) Notice of References Cited (PTO-892)	4) 🔲 Interview Su	mmary (PTO-413)	
 2) Notice of Draftsperson's Patent Drawing Review (PTO-948 3) Information Disclosure Statement(s) (PTO-1449 or PTO/SE Paper No(s)/Mail Date 	Paper No(s)	Mail Date commal Patent Application (PTO-152)	

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DETAILED ACTION

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-5, and 7-15, drawn to a method of alleviating drug resistant hypercalcemia using an antibody directed towards PTHrP.

Group II, claim(s) 1-6, and 14, drawn to a method of alleviating drug resistant hypercalcemia using an antagonist directed towards PTHrP receptor.

The inventions listed as Groups I-II do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: The inventions of Group I and II are different because they employ different mechanisms and different chemical entities of alleviating drug resistant hypercalcemia. The invention of Group I uses an antibody directed towards a ligand, while the invention of Group II uses an antagonist directed towards a receptor.

Claims 1-5 link(s) inventions I and II. The restriction requirement of the linked inventions is subject to the nonallowance of the linking claim(s), claims 1-5. Upon the allowance of the linking claim(s), the restriction requirement as to the linked inventions

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shall be withdrawn and any claim(s) depending from or otherwise including all the limitations of the allowable linking claim(s) will be entitled to examination in the instant application. Applicant(s) are advised that if any such claim(s) depending from or including all the limitations of the allowable linking claim(s) is/are presented in a continuation or divisional application, the claims of the continuation or divisional application may be subject to provisional statutory and/or nonstatutory double patenting rejections over the claims of the instant application. Where a restriction requirement is withdrawn, the provisions of 35 U.S.C. 121 are no longer applicable. *In re Ziegler*, 44 F.2d 1211, 1215, 170 USPQ 129, 131-32 (CCPA 1971). See also MPEP § 804.01.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **J. Jason Galvez**, **Ph.D**. whose telephone number is **571-272-2935**. The examiner can normally be reached Monday through Friday 9 AM to 5 PM. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, **Brenda Brumback**, **Ph.D**. can be reached at **571-272-0887**.

The fax phone number for the organization where this application or proceeding is assigned is **703-872-9306**. Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or

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11. The therapeutic agent according to claim 7, wherein the antibody is in a humanized form.

- 12. The therapeutic agent according to claim 11, wherein the humanized antibody is humanized #23-57-137-1 antibody.
- (Amended) The therapeutic agent according to any one of claims 1 or 2, wherein the drug-resistant hypercalcemia is caused by cancer.

RULE 1.121

1418. (New) The therapeutic agent according to claim 5, wherein the active ingredient is chosen from at least one of

- a) an antagonist for the PTHrP receptor;
- b) an anti-PTHrP antibody;
- c) a fragment of an anti-PTHrP antibody and/or a modified form of the fragment.
- 15. (New) The anti-PTHrP antibody of claim 14, wherein antibody is chosen from at least one of a human antibody, a humanized antibody, and a chimaric antibody.
- 16. (New) The therapeutic agent according to claim 5, wherein the drug-resistant hypercalcemia is caused by cancer.

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PATENT EXAMINER